Amendments to the Claims

- 1. (Currently amended) A composition for administration of a nucleic acid, comprising:
 - (a) liposomes comprised of
 - (i) a lipid having the formula

$$z \longrightarrow_{n} L \longrightarrow_{0} Q \xrightarrow{R^{1}} R^{2}$$

where each of R¹ and R² is an alkyl or alkenyl chain having between 8-24 carbon atoms, and each of R¹ and R² are independently selected;

$$n = [[1]]0-20;$$

L is selected from the group consisting of (1) -X-(C=O)-Y-[[CH₂-]], (2) -X-(C=O)-, and (3) -X-[[CH₂-]], where X and Y are independently selected from oxygen, NH and a direct bond;

Z is a weakly basic moiety that has a pK of less than 7.4 and greater than about 4.0; and

(ii) a compound having the general structure:

wherein R^3 is a hydrophilic polymer comprising a linkage for attachment to the dithiobenzyl moiety; R^4 is selected from the group consisting of H, alkyl and aryl; R^5 is selected from the group consisting of $O(C=O)R^7$, $S(C=O)R^7$, and $O(C=S)R^7$; R^7 comprises an amine-containing lipid; and R^6 is selected from the group consisting of H, alkyl and aryl; and where orientation of CH_2 - R^5 is selected from the ortho position and the para position; and

(b) a nucleic acid associated with said liposomes.

- 2. (Original) The composition of claim 1, wherein X is NH and Y is oxygen.
- 3. (Currently amended) The composition of claim 1, wherein L is a carbamate linkage, an ester linkage, or a carbonate linkage.
- 4. (Currently amended) The composition of claim 1, wherein L is NH-(C=O)-O[[-CH₂]].
 - 5. (Original) The composition of claim 1, wherein Z is an imidazole.
- 6. (Original) The composition of claim 1, comprising between 1-80 mole percent of the lipid.
- 7. (Original) The composition of claim 1, wherein Z is a moiety having a pK value between 5.0-6.5.
- 8. (Original) The composition of claim 1, wherein each of R¹ and R² is an unbranched alkyl or alkenyl chain having between 8-24 carbon atoms.
 - 9. (Original) The composition of claim 8, wherein each of R¹ and R² is C₁₇H₃₅.
 - 10. (Original) The composition of claim 1, wherein n is between 1-10.
- 11. (Original) The composition of claim 1, wherein R^6 is H and R^4 is selected from the group consisting of CH_3 , C_2H_5 and C_3H_8 .
- 12. (Original) The composition of claim 1, wherein the amine-containing lipid comprises either a single hydrocarbon tail or a double hydrocarbon tail.

- 13. (Original) The composition of claim 1, wherein the amine-containing lipid is a phospholipid having a double hydrocarbon tail.
 - 14. (Original) The composition of claim 1, wherein R⁴ and R⁶ are alkyls.
- 15. (Original) The composition of claim 1, wherein R³ is selected from the group consisting of polyvinylpyrrolidone, polyvinylmethylether, polymethyloxazoline, polyethyloxazoline, polyhydroxypropyloxazoline, polyhydroxypropyl-methacrylamide, polymethacrylamide, polyhydroxypropylmethacrylate, polyhydroxyethylacrylate, hydroxymethylcellulose, hydroxyethylcellulose, polyethyleneglycol, polyaspartamide, copolymers thereof, and polyethyleneoxide-polypropylene oxide.
 - 16. (Original) The composition of claim 1, wherein R³ is polyethyleneglycol.
- 17. (Original) The composition of claim 16, wherein R^6 is H and R^4 is CH_3 or C_2H_5 .
- 18. (Original) The composition of claim 1, wherein said liposomes include between 5-20 mole percent of the compound.
- 19. (Original) The composition of claim 1, further including a therapeutic compound entrapped in the liposomes.
- 20. (Original) The composition of claim 1, wherein said nucleic acid is entrapped in at least a portion of said liposomes.
- 21. (Original) The composition of claim 20, wherein the nucleic acid is selected from DNA, RNA, fragments thereof and oligonucleotides.

- 22. (Original) The composition of claim 1, further including a ligand for targeting the liposomes to a target site, said ligand covalently attached to a distal end of the hydrophilic polymer R³ on said compound.
- 23. (Original) The composition of claim 22, wherein the ligand has binding affinity for endothelial tumor cells for internalization by such cells.
- 24. (Original) The composition of claim 22, wherein the ligand is selected from the group consisting of E-selectin, Her-2 and FGF.
- 25. (Original) The composition of claim 22, wherein said ligand is selected from the group consisting of c-erbB-2 protein product of the HER2/neu oncogene, epidermal growth factor (EGF) receptor, basic fibroblast growth receptor (basic FGF) receptor, vascular endothelial growth factor receptor, E-selectin receptor, L-selectin receptor, P-selectin receptor, folate receptor, CD4 receptor, CD19 receptor, αβ integrin receptors, and chemokine receptors.
- 26. (Original) The composition of claim 1, wherein said liposomes further comprise a cationic lipid.